This listing of the claims will replace all prior versions and listings. Claims 1-7 are

currently pending in this application. Claim 1 has been amended.

**LISTING OF CLAIMS:** 

Claim 1 (currently amended): A method of assembling PCR fragments, comprising

a) making a first PCR fragment with first and second primers, wherein the second primer

comprises a modified nucleotide that can be removed by a DNA repair enzyme, resulting

in a 3' overhang, and wherein the first PCR fragment comprises a first site specific

recombinase site;

b) treating the first PCR fragment with a DNA repair enzyme to generate a 3' overhang and

immobilizing the first PCR fragment on a solid support or vice versa;

c) making a second PCR fragment with third and fourth primers, wherein the third and

fourth primers each comprises a modified nucleotide that can be removed by a DNA

repair enzyme resulting in a 3' overhang;

d) treating the second PCR fragment with a DNA repair enzyme to generate a 3' overhang;

e) annealing and ligating the first and second PCR fragments;

f) optionally repeating steps c, d and e until a last PCR fragment is added to the growing

chain to produce an assembled fragment, wherein the last PCR fragment comprises a

second site specific recombinase site; and

g) simultaneously removing and circularizing the assembled fragment from the solid support

with a site specific recombinase in a single step.

Claim 2 (original): The method of claim 1, where one of the PCR fragments comprises an origin

of replication and a selectable marker.

Claim 3 (original): The method of claim 1, wherein the first PCR fragment or the last PCR

fragment comprises an origin of replication and a selectable marker.

Page 2 of 4

App. No.: 10/699,511 Docket No.: 31175413-002002 (PATENT)

Response to Office Action mailed July 27, 2006

Claim 4 (original): The method of claim 1, wherein the site specific recombinase is CRE and the site specific recombinase site is lox.

- Claim 5 (original): The method of claim 1, wherein the nucleotide is deoxyuridine and the DNA repair enzyme is uracil-DNA-glycosylase followed by T<sub>4</sub> endonuclease V.
- Claim 6 (original): The method of claim 5, wherein the assembled DNA is greater than 30 kb.
- Claim 7 (original): The method of claim 5, wherein the assembled DNA is greater than 30, 40, 50, 75, 100, 125, 150, 200, 250, 300, 350, 400, 450, 500, 750, 1000 or 1500 kb.